

# Identifying factors associated with Covid-19 related deaths during the first wave of the pandemic in Europe

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## **Abstract**

Aim: To clarify the high variability in Covid-19-related deaths during the first wave of the pandemic, we conducted a modeling study using publicly available data.

Materials and Methods: We used 13 population- and country-specific variables to predict the number of population-standardized Covid-19-related deaths in 43 European countries using generalized linear models: percentage of test-standardized number of SARS-CoV-2-cases, population density, life expectancy, severity of governmental responses, influenza-vaccination coverage, vitamin D status, smoking and diabetes prevalence, cardiovascular disease death rate, number of hospital beds, gross domestic product, human development index and percentage of people older than 65 years.

Results: We found that test-standardized number of SARS-CoV-2-cases and flu vaccination coverage in the elderly were the most important predictors, together with vitamin D status, gross domestic product, population density and government response severity explaining roughly two-thirds of the variation in Covid-19 related deaths. The latter variable was positively, but only weakly associated with the outcome, i.e., deaths were higher in countries with more severe government response. Higher flu vaccination coverage and low vitamin D status were associated with more Covid-19 related deaths. Most other predictors appeared to be negligible.

Conclusion: Adequate vitamin D levels are important, while flu-vaccination in the elderly and stronger government response were putative aggravating factors of Covid-19 related deaths. These results may inform protection strategies against future infectious disease outbreaks.

## 1 Introduction

The SARS-Corona-Virus 2 (CoV2) pandemic caused an unprecedented worldwide public health crisis by its impact on basically every system of human organization (1,2). Untreated Covid-19 disease may lead to severe atypical pneumonia (3,4), a cytokine storm and other potentially lethal sequelae (5–7). Other potential factors, such as host factors or population factors, were not much considered in the scientific and political discourse. For example, there is now strong evidence that vitamin D status predicts the risk for and outcome of Covid-19 infections (8–13). We also know that demographics play a role, as initially mostly elderly patients with a mean age above 70 years have been severely affected (14,15). However, during the initial phase of the CoV2 outbreak, there was a wide variation in lethality across countries and regions. This variation is partially shrouded by the fact that most agencies and their dashboards propagate unstandardized figures of cases and deaths. A publication that estimated excess death rates in the US during the early time of the CoV2 pandemic as compared with the same months of previous years reveals a wide variation from -71,9 deaths per 100.000 inhabitants in North Dakota to 299,1 deaths per 100.000 inhabitants in New York City, with seven states actually exhibiting less excess mortality than in the previous comparison years, and 12 US states presenting with excess mortality figures below 10 per 100.000 inhabitants (16). The same is true for Europe: Miles and colleagues listed excess deaths of 21% for Spain, 20% for the UK, 18% for Italy down to 6% for Sweden, 3% for Portugal, -1% for Germany, -3% for Denmark and -4% for Norway during the first wave of the pandemic (17).

In order to be better prepared for future infectious disease outbreaks, there is clearly a need to understand what might have caused such variation in death numbers during the first wave of the pandemic. Are there population variables, public health variables, or individual-specific factors that can be identified that make this variation understandable? This was the guiding question of this modeling study.

## 2 Materials and Methods

We extracted data for 44 European countries for which the number of Covid-19 related deaths per 1.000.000 inhabitants up until 31<sup>st</sup> August 2020 was known. This date was chosen since it approximately marked the end of the first infection wave in Europe (18,19). The following 13 variables were used as putative predictors of the dependent variable “standardized Covid-19 related deaths” which we subsequently refer to as “y” (Supplementary Table 1): (i) the test-standardized number of cases (in %), calculated as the number of cases in a country divided by the number of tests in that country  $\times 100$ ; (ii) the influenza (flu) vaccination rate in the elderly; (iii) life expectancy (in years); (iv) the population density (people per km<sup>2</sup>); (v) mean Government Response Severity Index (GRSI) that describes the number and severity of non-pharmaceutical interventions employed between 15th March and 15th August 2020; (vi) vitamin D status (25(OH)D < 50nmol/l/  $\geq 50$ nmol/l); (vii) cardiovascular disease (CVD) death rate; (viii) diabetes prevalence; (ix) smoking habits (average percentage of male and female smokers); (x) percentage of elderly (people older than 65 years); (xi) gross domestic product; (xii) human development index; (xiii) hospital beds (number of beds per thousand inhabitants). The data sources are given in Supplementary Data File 1.

Because the distribution of  $y$  followed a gamma distribution well (Figure 1), we calculated generalized linear models (GLMs) on a gamma-distributed variable with a log-link function. Since a log-transformation produced an outcome variable with an approximately normal distribution (Shapiro-Wilk normality test  $p=0.864$ , Figure 1), we also calculated standard multiple linear regression models (LRMs) on  $\log(y)$ . During the initial check of modeling assumptions, one outlier (Andorra) was identified and removed from the sample (Supplementary Data File 1).

The final sample thus included 43 European countries of which 40 had known flu vaccination rates, 37 had known flu vaccination rates and GRSI values, and 31 had no missing variable values. To utilize as many cases as possible for multivariable modeling (20) missing variable values were imputed with multiple imputation by chained equations using the R package 'mice' (21). A total of 100 imputation data sets were created. Each was used to fit the regression models, and the model parameters were averaged over all 100 model fits.

Different regression models were pre-specified according to plausible scientific hypothetical explanations for Covid-19 related deaths, reflecting the scientific practice of evaluating multiple pre-specified working hypotheses for their ability to explain the observed data (22). To complement the set of pre-specified hypotheses, one additional model was built using the Least Absolute Shrinkage and Selection Operator (LASSO), a variable selection method that shrinks the regression coefficients of less important predictors to zero (23). Instead of performing multiple null hypothesis testing, we ranked the different models according to their evidence constituted by the data by using the bias-corrected Akaike Information criterion (22).

Because of the skewed distribution of some of the predictors, we first fitted a univariable model for each predictor and its log transform, and chose to use the latter for multivariable modeling if it resulted in a AICc reduction by at least 2 compared to its non-transformed values. In this way, it was found that test-standardized cases, population density and CVD death rate resulted in significantly better model fits as log-transformed variables.

As the simplest hypothesis, we assumed that the number of deaths could be predicted by the number of test-standardized cases:

$$y \sim \log(\text{test-standardized cases}) \quad (1)$$

The second most-plausible simple hypothesis was that in addition to the number of cases, the severity of governmental responses, whose concept was to prevent infections, would allow better predictions of the outcome:

$$y \sim \log(\text{test-standardized cases}) + \text{GRSI} \quad (2)$$

A third model was motivated by an interesting paper showing that the flu vaccination rate in the elderly was significantly correlated at  $r = .68$  with Covid-19 related deaths in Europe (24). Furthermore, early clinical data have indicated that vitamin D has protective effects against Covid-19, which would be expected based on its immune-modulatory functions (1). Finally, a modeling study by Liang et al. found that the number of hospital beds in a country was

associated with decreased Covid-19 mortality (25). In model 3, we therefore investigated the impact of flu vaccinates rates, nation-wide vitamin D status and the number of hospital beds, which all could be seen as features of the health care system:

$$y \sim \log(\text{test-standardized cases}) + \text{vitamin D status} + \text{hospital beds} + \text{flu vaccination rate} \quad (3)$$

During the first wave of the pandemic, elderly people were far more susceptible to Covid-19 related deaths (19,26), whereby an inverse association between vitamin D levels and Covid-19 severity was shown in an Italian study (27). These findings motivated the construction of a fourth model which attempted to predict Covid-19 related deaths based on vitamin D status and demographics:

$$y \sim \log(\text{test-standardized cases}) + \text{vitamin D status} + \text{life expectancy} + \text{percentage of elderly} \quad (4)$$

Besides old age, it was soon revealed during the early phase of the pandemic that cardiovascular disease (19), diabetes (28) and smoking (29,30) are associated with Covid-19 severity. The fifth hypothesis therefore assumed that population-level morbidity predictors would be relevant for predicting Covid-19 related deaths:

$$y \sim \log(\text{test-standardized cases}) + \text{vitamin D status} + \text{smoking habits} + \log(\text{CVD death rate}) + \text{diabetes prevalence} \quad (5)$$

Previous modeling studies also tested for associations between Covid-19 related deaths and different country-specific predictors such as the gross domestic product and percentage of the elderly (25,31). The sixth hypothesis therefore correlated Covid-19 deaths with such country-specific predictors:

$$y \sim \log(\text{test-standardized cases}) + \text{vitamin D status} + \log(\text{population density}) + \text{life expectancy} + \text{gross domestic product} + \text{human development index} + \text{percentage of elderly} \quad (6)$$

The seventh model was the full model using all 13 predictors which was included as a reference model (22).

Finally, an eighth model was built using a data-driven approach. To this aim, for each imputed dataset the most important variables were selected from the full set of 13 predictors using the LASSO method in a LRM predicting  $\log(y)$ . LASSO performs variable selection by shrinking the regression coefficients of the less important predictors to zero (32). The following variables were selected into the majority (>50) of models:

$$y \sim \log(\text{test-standardized cases}) + \text{vitamin D status} + \text{GRSI} + \text{flu vaccination rate} + \log(\text{CVD death rate}) + \log(\text{population density}) + \text{GDP} \quad (8)$$

The best model was identified as the one with the smallest AICc, and all other models were compared to the best model by computing AICc differences  $\Delta_i$ , probabilities  $w_i$  of model  $i$  being the best model (in the Kullback-Leibler information sense) and evidence ratios  $E_{i,j} =$

$w_i/w_j$  (22). Model adequacy was measured by  $R^2$ , the proportion of variance explained by the predictors; for the GLMs a Kullback-Leibler divergence-based  $R^2$  measure was used (33).

All analyses were calculated with R version 4.0.2, and statistical significance was defined as p-values  $<0.01$ . A detailed description of the statistics is given in Supplementary Data File 1.

### 3 Results

Figure 2 shows a so-called corrgram (34) for the 13 variables that we used for modeling, whereby only the significant ( $p < 0.01$ ) correlations have been depicted. Smoking prevalence and cardiovascular disease death rate, but not diabetes prevalence, were inversely and significantly correlated with life expectancy, the human development index and gross domestic product. No significant correlations existed for vitamin D levels with any of the other variables.

The results of both the GLMs (assuming a Gamma distribution for the outcome variable  $y$ ) and the LRMs fitted to  $\log(y)$  are presented in Table 1. The GLMs and LRMs yielded qualitatively similar results for all eight hypotheses considered. Test-standardized cases alone were able to explain about 20% of the variance in  $y$  or  $\log(y)$ , respectively, while the full model (model 7) was able to explain 64-67% as judged by the adjusted KL- $R^2$  values. As expected, test-standardized cases were positive and the most significant predictors of Covid-19 related deaths in all models. As also expected, sufficient vitamin D status was associated with fewer deaths, although the association was only significant at the conventional  $p = 0.05$  level in models 7 and 8, but not at  $p = 0.01$ , as stipulated. Surprisingly, however, it was found that the GRSI was no significant predictor of Covid-19 related deaths, and even exhibits a positive association (i.e. more stringent measures predicting higher death rates). Also surprisingly, flu vaccination rates were significantly and positively related to the outcome, i.e., there were more deaths in countries with higher flu-vaccination coverage. The number of hospital beds, percentage of elderly, human development index and smoking and diabetes prevalence, were not strongly associated with Covid-19 related deaths. In contrast, the GDP was found to be a significant predictor in models 7 and 8 with a higher GDP being associated with more Covid-19 related deaths.

A ranking of the different models is given in Table 2. The evidence clearly favored the data-driven model 8 which was built by first selecting variables using the LASSO method. Compared with this model, all other models were basically ruled out by the strength of evidence. This shows that a specific combination of government-, population- and country-specific factors were important for predicting Covid-19-related deaths. This final model 8 was thereby able to explain roughly two-thirds of the variance in outcomes, similar to the full model 7, but using seven less predictors.

In order to check if our results are dependent on the imputation of missing variables, we refitted the final GLM and LRM model to the original dataset with missing variables removed (Table 3). These models resulted in qualitatively very similar results as model 8 in Table 1 and confirmed that the two most important predictors of standardized COVID-19 deaths were

again flu vaccination rates and the number of test-standardized cases, which were both positively associated with the outcome. GDP was also confirmed as a significant and positively associated predictor of Covid-19 deaths, and vitamin D status now reached the threshold of statistical significance in the GLM ( $p=0.00573$ ).

#### 4 Discussion

Modeling Covid-19 related death rates in 43 European countries during the initial phase of the outbreak until August 2020, unravels some interesting findings:

- a) Unsurprisingly, test-standardized CoV2-cases predict the number of deaths. This variable on its own was able to explain about 20% of the variance.
- b) Surprisingly, the stringency of government responses correlated positively with Covid-19 related death rates, i.e., stricter government response was associated with more deaths; however, it was not a significant predictor.
- c) Also surprisingly, the second-most important predictor was the flu-vaccination coverage in the elderly: the higher this vaccination rate is, the more Covid-19 related deaths we see in a country.
- d) We confirmed that population-wide vitamin D status may have acted protectively against COVID-19 related deaths during the initial phase of the outbreak. It was a highly significant predictor in the best GLM fitted to the dataset with no missing variables (Table 3).
- e) Countries with a higher GDP experienced a higher Covid-19 associated death rate.

These findings are strengthened by the fact that two different models reached the same conclusions: a GLM predicting a gamma-distributed outcome variable with log-linked predictors and a standard multiple LRM with identity link functions of predictors on a log-transformed outcome variable.

It is easy to understand that more CoV2 cases translate into more Covid-19 related deaths. The importance of this predictor on its own is underlined by the fact that it is able to explain roughly 20% of Covid-19 related deaths. However, there remains variance to be explained by other factors. Although we do not assume we have captured all important variables, we have captured at least some as only six variables were able to explain about two-thirds of the total variance. A reassuring finding was that country-wide vitamin D status was inversely associated with Covid-19 related deaths, consistent with clinical and epidemiological data (8–13). Most surprising and most counterintuitive are the two findings that there are more Covid-19 related deaths in countries with higher flu vaccination coverage in the elderly, and, in addition, that the severity of governments' responses with non-pharmaceutical interventions was non-significant and counterintuitive in its effect (Table 1).

How can this strong association between flu vaccination rates and Covid-19 related deaths be explained? A careful randomized trial of flu vaccination in children showed that children who were vaccinated against influenza were better protected against influenza but suffered a fourfold higher risk of other respiratory virus dependent diseases (35). This might have to do with unknown mechanisms that disturb the ecology of pathogens, known as the virus

interference phenomenon. A study conducted during the 2017/2018 influenza season revealed that flu vaccination was associated with a 36% increased odds of contracting respiratory coronavirus diseases (odds ratio 95% confidence interval 1.14-1.63,  $p < 0.01$ ), while affording specific protection against influenza and parainfluenza viruses (36).

Thus, the negative impact of flu vaccination might have to do with several mechanisms: First, the virus interference phenomenon as shown for non-CoV2 coronaviruses (36); second, the fact that the immunological load on an organism that has to deal with a flu vaccine binds resources that cannot be mustered against a new and dangerous pathogen like CoV2. Third, it might also be the case that immune-enhancers in vaccines, such as aluminum derivatives which are potentially toxic, burden the organism and hamper natural immunity. For example, it was shown experimentally in chicks that aluminum can disturb vitamin D metabolism (37). Furthermore, it has been argued that influenza vaccines are produced in eggs and other cell-systems that are not routinely tested against corona-viruses. Hence, corona-virus proteins from other corona-viruses might be present in these vaccines and induce allergic reactions against the novel CoV2 (38). Although these biological mechanisms would support the hypothesis that higher flu vaccination rates increased Covid-19 mortality rates, we cannot rule out the possibility that flu vaccination rate is simply a non-causal confounder strongly associated with some other (untested) variable, so that further research is needed to resolve this issue. Our finding is also in contrast to data from the US (39,40). However, the correlation between flu vaccination and COVID-19 death rate in the US is much lower than in Europe (24), probably because there is little variation in influenza vaccine coverage in the US. Our results are derived from population-level data in Europe in the elderly, which might describe a specifically susceptible fraction of the population.

Non-pharmaceutical interventions were widely hailed in modeling studies as having prevented higher incidence figures of cases and deaths (41–43). While this might be true for some countries and some single interventions, some authors are skeptical (44–49). Careful modeling studies for Germany, for instance, show that, although Germany was comparatively early to react – first measures were introduced on March 8 and shortly after this a full country lockdown was enacted – the peak of the infection and of the reproduction numbers was reached in nearly all 420 German districts on or around March 8 and thus none of the non-pharmaceutical interventions could have been causally related to the reduction of cases, and hence deaths (50,51). The ensuing reduction of cases is a misattribution: it is not due to the lockdown, but obviously to the fact that the virus followed its own dynamic which needs to be better understood (52). Thus, there is independent evidence that non-pharmaceutical interventions are less effective than often thought. This would explain the weak association with Covid-19 related deaths in our analysis. Interestingly, our observation that the GRSI was positively associated with Covid-19 related deaths during the first phase of the pandemic replicates an earlier modeling study by Annaka which included data from 108 countries and in which this association was even significant in Ordinary Least Squares regression (31).

We find it quite remarkable that only six variables help to explain roughly two-thirds of the variation in Covid-19 related deaths. Because vitamin D status was one of them, it might be interesting to study other variables related to health. Vitamin D entered the best model



number 8 with a comparatively large regression coefficient and was highly significant in a GLM fit to the complete dataset. Vitamin D seems to be an important predictor, as models without it are clearly inferior. For example, removing vitamin D status as a predictor from the GLM 8 fitted to the complete dataset resulted in a significantly worse model fit (AICc = 424.4 vs. 362.0) and less efficiency in explaining variance ( $KL-R^2_{adj} = 0.684$  vs.  $KL-R^2_{adj} = 0.726$ ). Thus, as a theoretically and numerically strong predictor, vitamin D strongly improves model fit and therefore we conclude that vitamin D was protective against death during the first wave of the Covid-19 pandemic. Its lack of strong statistical significance in models fit to the imputed dataset is likely due to the coarse grained nature of our data and uncertainty in imputation of missing values.

The limitations of our approach need to be kept in mind:

First, there was collinearity among some of the variables we used for modeling (Figure 2). However, no significant correlations existed among the six variables in the best model (Figure 3). This was supported by the computation of variance inflation factors which were all  $<1.65$ , showing that there was no significant collinearity between these six variables. In particular, flu vaccination rate had the least collinearity with the other predictors (variance inflation factor = 1.2).

Second, we were unable to find flu vaccination data, GRSI and some other data for all countries. We tried to overcome this limitation through multiple imputation by chained equations, and the results were consistent with an analysis using only the cases for which every variable value was known.

Third, one potential problem we cannot remedy is the notorious unreliability of data or differences in the definition of cases, of deaths, and in reporting standards. This can be seen in the fact that Belgium is a clear outlier in all analyses that decreases the fit of the model. It is well known that the definition of Covid-19 related deaths in Belgium was more lenient than in other countries. Also, there is some evidence that some authoritarian governments tended to manipulate (downplayed) their Covid-19 death data (31), which could have biased our results. In his country-level model study, Annaka (31) accounted for such a putative reporting bias by including the HRV transparency index developed by Hollyer et al. (53) which he used as a proxy for data transparency. However, the HRV index was originally not intended for assessing the transparency of pandemic death reporting; in addition, its latest version (the one used by Annaka) dates to the year 2010 and was only available for 21 of the 43 countries included in our analysis (median index 4.403, range -0.685–5.636); that Denmark scored worst with an HRV index of only -0.685 also appears counterintuitive and questions the application of this index to judge the transparency of Covid-19 related deaths reporting. The fact that we restricted the analysis to European countries of which the large majority nowadays is characterized as democratic would have mitigated the putative effects of data transparency bias.

Vitamin D estimates also have several uncertainties, such as having been measured in rather small cohorts, in different years and during different times of the year. Whenever possible, we preferred vitamin D values from the literature that had been measured in elderly people and

during winter/spring. There was a weak positive correlation between a country's representative 25(OH)D concentrations and latitude (Kendall's  $\tau=0.255$ ,  $p=0.0438$ ), pointing towards vitamin D supplementation having a stronger influence on vitamin D status than living in southern latitudes. Also, Covid-19 reporting systems might be less reliable in some countries compared with others. These are the limits of our data and our analyses. But considering the fact that the whole world, politicians and public health officials use exactly the same data for their decisions should allow us to use them for analysis. One should remember that being a case, when considering the number of tests in a country, has only a weak relationship with becoming a fatality. It has been shown that the case fatality rate during the first wave was much less than previously assumed and estimated to be 0.15% (54). In Germany the case-fatality rate has been calculated from well documented cohorts to be 0.12 to 0.35% (55,56). The still widely circulating higher case fatality rates are due to the fact that they are largely calculated using raw, absolute figures without knowledge of the real prevalence (57). But also standardized figures might be unreliable. Often the same person is tested multiple times. Thus, we likely overestimate the number of cases by some margin. This would mean: the true link between being a case and becoming a fatality is probably even weaker.

Considering all these weaknesses our paper also has some strengths. First, care was taken to ensure that the essential requirements for linear modeling were met. Second, we pre-specified plausible hypotheses (expressed as GLMs or LRMs) and used a robust model comparison framework based on Kullback-Leibler information to compare them, in this way automatically incorporating penalties for potential overfitting. Third, restricting the analysis to Europe means that we have a comparatively homogeneous sample which nevertheless has enough variability. While all countries issued warnings the way NPIs were implemented differed widely, from suggestions and recommendations in Sweden to very strict stay-at-home orders that were policed in Spain, from nearly no regard in Belarus to strict political measures in Italy. Thus, we likely have seen a representative laboratory for the world, except that we do not cover any variance in ethnicity.

## **5 Conclusions**

In conclusion we see that Covid-19 related deaths during the first wave were most importantly dependent on the percentage of test-positive cases and flu-vaccination rate among the elderly in a country, whereby larger flu vaccination rates were associated with higher Covid-19 related deaths. The third important predictor was the GDP, followed by country-wide vitamin D status in the elderly, for which a causal relationship appears well supported by clinical and mechanistic evidence. These variables predict the variability in Covid-19 related deaths much better than the severity of governmental responses, the availability of hospital beds, smoking and diabetes prevalence or CVD death rates. Overall, we were able to show that a specific combination of government response-, population- and country-specific predictors was able to explain roughly two-thirds of the variance in Covid-19 related deaths. This might encourage others to look for additional factors that may explain the remainder of the variability in cases and deaths during the initial phases of the CoV2 outbreak. Ultimately, using the insights from modeling studies such as ours may help to be better prepared against future infectious disease outbreaks.

## **Conflict of Interest**

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

## **Author Contributions**

Conceptualization: HW; Methodology: RJK, HW; Formal analysis: RJK; Investigation: RJK, HW; Data Curation: RJK, HW; Writing - Original Draft: RJK, HW; Writing – Review & Editing: RJK, HW

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## **Ethical approval**

This study was a secondary analysis on publicly available data and hence no ethical approval was required.

## **Data availability**

The underlying data for this analysis are given in the Supplementary Data file 1

## **References**

1. Klement RJ. Systems Thinking About SARS-CoV-2. *Front Public Heal* (2020) **8**:585229. doi:10.3389/fpubh.2020.585229
2. Strelkovskii N, Rovenskaya E. Causal Loop Diagramming of Socioeconomic Impacts of COVID-19: State-of-the-Art, Gaps and Good Practices. *Systems* (2021) **9**:65.
3. Wu C, Chen X, Cai Y, Xia J, Zhou X, Xu S, Huang H, Zhang L, Zhou X, Du C, et al. Risk Factors Associated with Acute Respiratory Distress Syndrome and Death in Patients with Coronavirus Disease 2019 Pneumonia in Wuhan, China. *JAMA Intern Med* (2020) **180**:934–943. doi:10.1001/jamainternmed.2020.0994
4. Adhikari SP, Meng S, Wu Y, Mao Y-P, Ye R-X, Wang Q-Z, Sun C, Sylvia S, Rozelle S, Raat H, et al. Epidemiology, causes, clinical manifestation and diagnosis, prevention and control of coronavirus disease (COVID-19) during the early outbreak period: a scoping review. *Infect Dis Poverty* (2020) **9**:29. doi:10.21203/rs.2.24474/v1
5. Qin C, Zhou L, Hu Z, Zhang S, Yang S, Tao Y, Xie C, Ma K, Shang K, Wang W, et al. Dysregulation of immune response in patients with coronavirus 2019 (COVID-19) in Wuhan, China. *Clin Infect Dis* (2020) **71**:762–768. doi:10.1093/cid/ciaa248
6. Shi S, Qin M, Shen B, Cai Y, Liu T, Yang F, Gong W, Liu X, Liang J, Zhao Q, et al. Association of Cardiac Injury with Mortality in Hospitalized Patients with COVID-19

- in Wuhan, China. *JAMA Cardiol* (2020) **5**:802–810.  
doi:10.1001/jamacardio.2020.0950
7. Richardson S, Hirsch JS, Narasimhan M, Crawford JM, McGinn T, Davidson KW, Barnaby DP, Becker LB, Chelico JD, Cohen SL, et al. Presenting Characteristics, Comorbidities, and Outcomes among 5700 Patients Hospitalized with COVID-19 in the New York City Area. *JAMA - J Am Med Assoc* (2020) **323**:2052–2059.  
doi:10.1001/jama.2020.6775
  8. Singh S, Kaur R, Singh RK. Revisiting the role of vitamin D levels in the prevention of COVID-19 infection and mortality in European countries post infections peak. *Aging Clin Exp Res* (2020) **32**:1609–1612. doi:10.1007/s40520-020-01619-8
  9. Rhodes JM, Subramanian S, Laird E, Griffin G, Kenny RA. Perspective: Vitamin D deficiency and COVID-19 severity – plausibly linked by latitude, ethnicity, impacts on cytokines, ACE2 and thrombosis. *J Intern Med* (2021) **289**:97–115.  
doi:10.1111/joim.13149
  10. Radujkovic A, Hippchen T, Tiwari-Heckler S, Dreher S, Boxberger M, Merle U. Vitamin D deficiency and outcome of COVID-19 patients. *Nutrients* (2020) **12**:2757.  
doi:10.3390/nu12092757
  11. Entrenas Castillo M, Entrenas Costa LM, Vaquero Barrios JM, Alcalá Díaz JF, Miranda JL, Bouillon R, Quesada Gomez JM. “Effect of Calcifediol Treatment and best Available Therapy versus best Available Therapy on Intensive Care Unit Admission and Mortality Among Patients Hospitalized for COVID-19: A Pilot Randomized Clinical study”. *J Steroid Biochem Mol Biol* (2020) **203**:105751.  
doi:10.1016/j.jsbmb.2020.105751
  12. Annweiler G, Corvaisier M, Gautier J, Dubée V, Legrand E, Sacco G, Annweiler C. Vitamin D Supplementation Associated to Better Survival in Hospitalized Frail Elderly COVID-19 Patients: The GERIA-COVID Quasi-Experimental Study. *Nutrients* (2020) **12**:3377. doi:10.3390/nu12113377
  13. Maghbooli Z, Sahraian MA, Ebrahimi M, Pazoki M, Kafan S, Tabriz HM, Hadadi A, Montazeri M, Nasiri M, Shirvani A, et al. Vitamin D sufficiency, a serum 25-hydroxyvitamin D at least 30 ng/mL reduced risk for adverse clinical outcomes in patients with COVID-19 infection. *PLoS One* (2020) **15**:e0239799.  
doi:10.1371/journal.pone.0239799
  14. Grasselli G, Zangrillo A, Zanella A, Antonelli M, Cabrini L, Castelli A, Cereda D, Coluccello A, Foti G, Fumagalli R, et al. Baseline Characteristics and Outcomes of 1591 Patients Infected with SARS-CoV-2 Admitted to ICUs of the Lombardy Region, Italy. *JAMA - J Am Med Assoc* (2020) **323**:1574–1581. doi:10.1001/jama.2020.5394
  15. Onder G, Rezza G, Brusaferro S. Case-Fatality Rate and Characteristics of Patients Dying in Relation to COVID-19 in Italy. *JAMA - J Am Med Assoc* (2020) **323**:1775–1776. doi:10.1001/jama.2020.4683
  16. Weinberger DM, Chen J, Cohen T, Crawford FW, Mostashari F, Olson D, Pitzer VE, Reich NG, Russi M, Simonsen L, et al. Estimation of Excess Deaths Associated with the COVID-19 Pandemic in the United States, March to May 2020. *JAMA Intern Med* (2020) **180**:1336–1344. doi:10.1001/jamainternmed.2020.3391

17. Miles D, Stedman M, Heald A. Living with COVID-19: Balancing Costs against Benefits in the Face of the Virus. *Natl Inst Econ Rev* (2020) **253**:R60–R76. doi:10.1017/nie.2020.30
18. Di Gennaro F, Gualano G, Timelli L, Vittozzi P, Di Bari V, Libertone R, Cerva C, Pinnarelli L, Nisii C, Ianniello S, et al. Increase in tuberculosis diagnostic delay during first wave of the covid-19 pandemic: Data from an Italian infectious disease referral hospital. *Antibiotics* (2021) **10**:272. doi:10.3390/antibiotics10030272
19. Maximiano Sousa F, Roelens M, Fricker B, Thiabaud A, Iten A, Cusini A, Flury D, Buettcher M, Zukol F, Balmelli C, et al. Risk factors for severe outcomes for COVID-19 patients hospitalised in Switzerland during the first pandemic wave, February to August 2020: prospective observational cohort study. *Swiss Med Wkly* (2021) **151**:w20547. doi:10.4414/smw.2021.20547
20. Sterne JAC, White IR, Carlin JB, Spratt M, Royston P, Kenward MG, Wood AM, Carpenter JR. Multiple imputation for missing data in epidemiological and clinical research: potential and pitfalls. *BMJ Br Med J* (2009) **338**:b2393. doi:10.1136/bmj.b2393
21. Van Buuren S, Groothuis-Oudshoorn K. mice: Multivariate Imputation by Chained Equations in R. *J Stat Softw* (2011) **45**:1–67. doi:10.1177/0962280206074463
22. Anderson DR. *Model Based Inferences in the Life Sciences: A primer on Evidence*. 1st ed. New York: Springer Science+Business Media, LLC (2008).
23. James G, Witten D, Hastie T, Tibshirani R. *An Introduction to Statistical Learning with Applications in R*. New York: Springer Science+Business Media (2013).
24. EBMPHET Consortium. COVID-19 Severity in Europe and the USA: Could the Seasonal Influenza Vaccination Play a Role? [Preprint]. *SSRN* (2020) doi:10.2139/ssrn.3621446
25. Liang LL, Tseng CH, Ho HJ, Wu CY. Covid-19 mortality is negatively associated with test number and government effectiveness. *Sci Rep* (2020) **10**:12567. doi:10.1038/s41598-020-68862-x
26. Ioannidis JPA, Axfors C, Contopoulos-Ioannidis DG. Population-level COVID-19 mortality risk for non-elderly individuals overall and for non-elderly individuals without underlying diseases in pandemic epicenters. *Environ Res* (2020) **188**:109890. doi:10.1016/j.envres.2020.109890
27. Sulli A, Gotelli E, Casabella A, Paolino S, Pizzorni C, Alessandri E, Grosso M, Ferone D, Smith V, Cutolo M. Vitamin D and Lung Outcomes in Elderly Covid-19 Patients. *Nutrients* (2021) **13**:717. doi:10.3390/nu13030717
28. Peric S, Stulnig TM. Diabetes and COVID-19: Disease—Management—People. *Wien Klin Wochenschr* (2020) **132**:356–361. doi:10.1007/s00508-020-01672-3
29. Vardavas CI, Nikitara K. COVID-19 and smoking: A systematic review of the evidence. *Tob Induc Dis* (2020) **18**:20. doi:10.18332/tid/119324
30. Kashyap VK, Dhasmana A, Massey A, Kotnala S, Zafar N, Jaggi M, Yallapu MM,

- Chauhan SC. Smoking and COVID-19: Adding fuel to the flame. *Int J Mol Sci* (2020) **21**:6581. doi:10.3390/ijms21186581
31. Annaka S. Political regime, data transparency, and COVID-19 death cases. *SSM - Popul Heal* (2021) **15**:100832. doi:10.1016/j.ssmph.2021.100832
  32. Tibshirani R. Regression Shrinkage and Selection via the Lasso. *J R Stat Soc Ser B* (1996) **58**:267–288. doi:https://doi.org/10.1111/j.2517-6161.1996.tb02080.x
  33. Colin Cameron A, Windmeijer FAG. An R-squared measure of goodness of fit for some common nonlinear regression models. *J Econom* (1997) **77**:329–342. doi:https://doi.org/10.1016/S0304-4076(96)01818-0
  34. Friendly M. Corrgrams: Exploratory Displays for Correlation Matrices. *Am Stat* (2002) **56**:316–324. doi:10.1198/000313002533
  35. Cowling BJ, Fang VJ, Nishiura H, Chan K, Ng S, Ip DKM, Chiu SS, Leung GM, Peiris JSM. Increased Risk of Noninfluenza Respiratory Virus Infections Associated With Receipt of Inactivated Influenza Vaccine. *Clin Infect Dis* (2012) **54**:1778–1783. doi:10.1093/cid/cis307
  36. Wolff GG. Influenza vaccination and respiratory virus interference among Department of Defense personnel during the 2017–2018 influenza season. *Vaccine* (2020) **38**:350–354. doi:10.1016/j.vaccine.2019.10.005
  37. Henry HL, Norman AW. Interactions Between Aluminum and the Actions and Metabolism of Vitamin D3 in the Chick. *Calcif Tissue Int* (1985) **37**:484–490. doi:10.1007/BF02557831
  38. Arumugham V. Proteins that contaminate influenza vaccines have high homology to SARS-CoV-2 proteins thus increasing risk of severe COVID-19 disease and mortality. *Zenodo* (2020)
  39. Zanettini C, Omar M, Dinalankara W, Imada EL, Colantuoni E, Parmigiani G, Marchionni L. Influenza Vaccination and COVID19 Mortality in the USA. *medRxiv* (2020)2020.06.24.20129817. doi:10.1101/2020.06.24.20129817
  40. Zein JG, Whelan G, Erzurum SC. Safety of influenza vaccine during COVID-19. *J Clin Transl Sci* (2020) **5**:e49. doi:10.1017/cts.2020.543
  41. Flaxman S, Mishra S, Gandy A, Unwin HJT, Mellan TA, Coupland H, Whittaker C, Zhu H, Berah T, Eaton JW, et al. Estimating the effects of non-pharmaceutical interventions on COVID-19 in Europe. *Nature* (2020) **584**:257–261. doi:10.1038/s41586-020-2405-7
  42. Dehning J, Zierenberg J, Spitzner FP, Wibral M, Neto JP, Wilczek M, Priesemann V. Inferring change points in the spread of COVID-19 reveals the effectiveness of interventions. *Science* (2020) **369**:eabb9789. doi:10.1126/science.abb9789
  43. Hunter PR, Colón-González FJ, Brainard J, Rushton S. Impact of non-pharmaceutical interventions against COVID-19 in Europe: A quasi-experimental study. *medRxiv* (2020) doi:10.1101/2020.05.01.20088260

44. Klement RJ. The SARS-CoV-2 Crisis: Has Medicine Finally Entered a Reductionist Era? *Complement Med Res* (2020) **27**:207–208. doi:10.1159/000510453
45. Daunizeau J, Moran R, Mattout J, Friston K. On the reliability of model-based predictions in the context of the current COVID epidemic event: Impact of outbreak peak phase and data paucity. *medRxiv* (2020) doi:10.1101/2020.04.24.20078485
46. Moran RJ, Fagerholm ED, Cullen M, Daunizeau J, Richardson MP, Williams S, Turkheimer F, Leech R, Friston KJ. Estimating required ‘lockdown’ cycles before immunity to SARS-CoV-2: Model-based analyses of susceptible population sizes, ‘S0’, in seven European countries including the UK and Ireland [version 1; peer review: 1 approved with reservations, 1 not approved. *Wellcome Open Res* (2020) **5**:85. doi:10.1101/2020.04.10.20060426
47. Friston KJ, Parr T, Zeidman P, Razi A, Flandin G, Daunizeau J, Hulme OJ, Billig AJ, Litvak V, Price CJ, et al. Second waves, social distancing, and the spread of COVID-19 across America [version 1; peer review: awaiting peer review]. *Wellcome Open Res* (2021) **5**:103. Available at:  
<https://search.proquest.com/docview/2407844071?accountid=26724>  
[http://sfx.library.cdc.gov/cdc/?url\\_ver=Z39.88-2004&rft\\_val\\_fmt=info:ofi/fmt:kev:mtx:journal&genre=article&sid=ProQ:ProQ%3Ahalthcompleteshell&atitle=Second+waves%2C+social+distancing%2C+and+t](http://sfx.library.cdc.gov/cdc/?url_ver=Z39.88-2004&rft_val_fmt=info:ofi/fmt:kev:mtx:journal&genre=article&sid=ProQ:ProQ%3Ahalthcompleteshell&atitle=Second+waves%2C+social+distancing%2C+and+t)
48. Chin V, Ioannidis JPA, Tanner MA, Cripps S. Effect Estimates of COVID-19 Non-Pharmaceutical Interventions are Non-Robust and Highly Model-Dependent. *J Clin Epidemiol* (2021) doi:10.1016/j.jclinepi.2021.03.014
49. Kuhbandner C, Homburg S. Commentary: Estimating the effects of non-pharmaceutical interventions on COVID-19 in Europe. *Front Med* (2020) **7**:580361. doi:10.3389/fmed.2020.580361
50. Wieland T. A phenomenological approach to assessing the effectiveness of COVID-19 related nonpharmaceutical interventions in Germany. *Saf Sci* (2020) **131**:104924. doi:10.1016/j.ssci.2020.104924
51. Wieland T. Flatten the curve! modeling SARS-CoV-2/COVID-19 growth in Germany at the county level. *REGION* (2020) **7**:43–83. doi:10.18335/REGION.V7I2.324
52. Kuhbandner C, Homburg S, Walach H, Hockertz S. Was Germany’s Lockdown in Spring 2020 Necessary? How bad data quality can turn a simulation into a dissimulation that shapes the future. *Futures* (2022) **135**:102879. doi:10.1016/j.futures.2021.102879
53. Hollyer JR, Peter Rosendorff B, Vreeland JR. Measuring transparency. *Polit Anal* (2014) **22**:413–434. doi:10.1093/pan/mpu001
54. Ioannidis JPA. Reconciling estimates of global spread and infection fatality rates of COVID-19: an overview of systematic evaluations. *Eur J Clin Invest* (2021) doi:10.1111/eci.13554
55. Streeck H, Schulte B, Kümmerer BM, Richter E, Höller T, Fuhrmann C, Bartok E, Dolscheid-Pommerich R, Berger M, Wessendorf L, et al. Infection fatality rate of

SARS-CoV2 in a super-spreading event in Germany. *Nat Commun* (2020) **11**:1–12.  
doi:10.1038/s41467-020-19509-y

56. Schrappe M, François-Kettner H, Gruhl M, Hart D, Knieps F, Manow P, Pfaff H, Püschel K, Glaeske G. Die Pandemie durch SARS-CoV-2/Covid-19 – der Übergang zur chronischen Phase. *Monit Versorgungsforsch* (2020) **13**:35–68.  
doi:10.24945/MVF.05.20.1866-0533.2248
57. Klement RJ, Bandyopadhyay PS. The Epistemology of a Positive SARS-CoV-2 Test. *Acta Biotheor* (2020) doi:10.1007/s10441-020-09393-w

### Figure captions

**Figure 1:** Left: Observed distribution of the outcome variable “Covid-19 related deaths per 1 million inhabitants” and a gamma distribution (rate = 0.00438, shape = 0.9270) fitted through maximum likelihood estimation. Right: Observed distribution of the log-transformed outcome variable, with a best-fit normal distribution.

Figure 2: Corrgram showing the Spearman correlation coefficients for all significant correlations among the 13 variables used for modeling (N=31, countries with missing variables removed).

Figure 3: Corrgram showing the Spearman correlation coefficients for all pairs of the six variables included in the best model (N=31, countries with missing variables removed). The strongest correlation was observed between vitamin D and the gross domestic product which was almost significant (p=0.013).



Table 1: Parameters of the generalized linear models fitted to standardized deaths and and linear models fitted to the logarithm of standardized deaths

<b>Model 1</b>	<b>Generalized linear model</b>				<b>Linear model</b>			
<b>Variable</b>	<b>Regression coefficient</b>	<b>p-value</b>	<b>KL-R<sup>2</sup></b>	<b>KL-R<sup>2</sup><sub>adj.</sub></b>	<b>Regression coefficient</b>	<b>p-value</b>	<b>R<sup>2</sup></b>	<b>R<sup>2</sup><sub>adj.</sub></b>
log(test-standardized cases [%])	0.600 (0.164)	0.00076	0.218	0.199	0.568 (0.165)	0.0014*	0.225	0.206
<b>Model 2</b>	<b>Generalized linear model</b>				<b>Linear model</b>			
<b>Variable</b>	<b>Regression coefficient</b>	<b>p-value</b>	<b>KL-R<sup>2</sup></b>	<b>KL-R<sup>2</sup><sub>adj.</sub></b>	<b>Regression coefficient</b>	<b>p-value</b>	<b>R<sup>2</sup></b>	<b>R<sup>2</sup><sub>adj.</sub></b>
log(test-standardized cases [%])	0.609 (0.159)	0.00047	0.297	0.261	0.498 (0.165)	0.0045*	0.286	0.250
GRSI	0.037 (0.017)	0.035			0.032 (0.017)	0.075		
<b>Model 3</b>	<b>Generalized linear model</b>				<b>Linear model</b>			
<b>Variable</b>	<b>Regression coefficient</b>	<b>p-value</b>	<b>KL-R<sup>2</sup></b>	<b>KL-R<sup>2</sup><sub>adj.</sub></b>	<b>Regression coefficient</b>	<b>p-value</b>	<b>R<sup>2</sup></b>	<b>R<sup>2</sup><sub>adj.</sub></b>
log(test-standardized cases [%])	0.729 (0.145)	1.9×10 <sup>-5*</sup>	0.524	0.474	0.678 (0.157)	0.00013*	0.491	0.437
Vitamin D status (sufficient vs. deficient)	-0.299 (0.367)	0.425			-0.510 (0.348)	0.154		
Hospital beds per 1000	0.025 (0.068)	0.710			0.011 (0.064)	0.859		
Flu vaccination rate [%]	0.032 (0.007)	3.1×10 <sup>-5*</sup>			0.029 (0.007)	0.00029*		
<b>Model 4</b>	<b>Generalized linear model</b>				<b>Linear model</b>			
<b>Variable</b>	<b>Regression coefficient</b>	<b>p-value</b>	<b>KL-R<sup>2</sup></b>	<b>KL-R<sup>2</sup><sub>adj.</sub></b>	<b>Regression coefficient</b>	<b>p-value</b>	<b>R<sup>2</sup></b>	<b>R<sup>2</sup><sub>adj.</sub></b>
log(test-standardized cases [%])	0.730 (0.131)	2.9×10 <sup>-6*</sup>	0.498	0.446	0.695 (0.164)	0.00016*	0.424	0.363
Vitamin D status (sufficient vs. deficient)	-0.472 (0.287)	0.109			-0.416 (0.374)	0.275		
Life expectancy [years]	0.189 (0.040)	4.1×10 <sup>-5*</sup>			0.155 (0.050)	0.0043*		
Elderly [%]	-0.033 (0.051)	0.533			0.005 (0.069)	0.943		
<b>Model 5</b>	<b>Generalized linear model</b>				<b>Linear model</b>			
<b>Variable</b>	<b>Regression coefficient</b>	<b>p-value</b>	<b>KL-R<sup>2</sup></b>	<b>KL-R<sup>2</sup><sub>adj.</sub></b>	<b>Regression coefficient</b>	<b>p-value</b>	<b>R<sup>2</sup></b>	<b>R<sup>2</sup><sub>adj.</sub></b>
log(test-standardized cases [%])	0.709 (0.151)	8.3×10 <sup>-5*</sup>	0.551	0.490	0.735 (0.158)	5.3×10 <sup>-5*</sup>	0.504	0.437
Vitamin D status (sufficient vs. deficient)	-0.363 (0.307)	0.248			-0.488 (0.346)	0.170		

deficient)								
Smoking prevalence [%]	-0.007 (0.029)	0.806			-0.006 (0.030)	0.844		
log(CVD death rate)	-0.999 (0.388)	0.020			-0.985 (0.392)	0.020		
Diabetes prevalence [%]	-0.09 (0.07)	0.185			-0.112 (0.080)	0.174		
<b>Model 6</b>								
log(test-standardized cases [%])	0.961 (0.171)	6.0×10 <sup>-6</sup> *	0.557	0.484	0.894 (0.198)	0.00013*	0.516	0.436
log(population density [km <sup>-2</sup> ])	0.222 (0.113)	0.058			0.127 (0.133)	0.325		
Life expectancy [years]	-0.011 (0.076)	0.883			0.011 (0.087)	0.886		
Gross domestic product	2.6×10 <sup>-5</sup> (1.5×10 <sup>-5</sup> )	0.088			3.8×10 <sup>-5</sup> (1.7×10 <sup>-5</sup> )	0.037		
Human development index	1.94 (5.52)	0.726			-3.9 (6.4)	0.549		
Elderly [%]	0.077 (0.062)	0.224			0.110 (0.074)	0.152		
<b>Model 7</b>								
log(test-standardized cases [%])	0.968 (0.148)	1.7×10 <sup>-6</sup> *	0.773	0.671	0.951 (0.172)	1.3×10 <sup>-5</sup> *	0.756	0.647
GRSI	0.030 (0.015)	0.055			0.030 (0.018)	0.118		
Vitamin D status (sufficient vs. deficient)	-0.716 (0.326)	0.043			-0.763 (0.372)	0.055		
Flu vaccination rate	0.019 (0.006)	0.0055*			0.020 (0.007)	0.011		
Life expectancy [years]	-0.046 (0.112)	0.685			-0.070 (0.123)	0.575		
log(population density [km <sup>-2</sup> ])	0.092 (0.107)	0.399			0.062 (0.124)	0.623		
Smoking prevalence [%]	-0.014 (0.022)	0.520			-0.022 (0.025)	0.350		
log(CVD death rate)	-0.037 (0.769)	0.962			-0.165 (0.936)	0.862		
Diabetes prevalence [%]	-0.058 (0.069)	0.414			-0.047 (0.082)	0.572		
Hospital beds per 1000	0.032 (0.072)	0.657			0.065 (0.082)	0.433		
GDP	3.8×10 <sup>-5</sup> (1.2×10 <sup>-5</sup> )	0.0055*			4.2×10 <sup>-5</sup> (1.4×10 <sup>-5</sup> )	0.0074*		
Human development index	-2.5 (5.2)	0.630			-2.8 (6.2)	0.654		
Elderly [%]	0.088 (0.057)	0.143			0.097 (0.065)	0.150		
<b>Model 8</b>								
log(test-standardized cases [%])	0.844 (0.123)	9.1×10 <sup>-7</sup> *	0.727	0.681	0.809 (0.147)	1.1×10 <sup>-5</sup> *	0.689	0.634
GRSI	0.021 (0.011)	0.071			0.021 (0.014)	0.154		
Vitamin D status (sufficient vs.	-0.704 (0.283)	0.023			-0.801 (0.332)	0.024		

deficient)							
Flu vaccination rate [%]	0.021 (0.005)	0.00021*			0.021 (0.006)	0.0015*	
log(population density [km <sup>-2</sup> ])	0.127 (0.097)	0.209			0.099 (0.111)	0.379	
GDP	2.8×10 <sup>-5</sup> (7.3×10 <sup>-6</sup> )	0.0070*			3.0×10 <sup>-5</sup> (9.0×10 <sup>-6</sup> )	0.0023*	

GDP: Gross domestic product; GRSI: Government Response Severity Index; CVD: Cardiovascular disease; \*p<0.01 (statistically significant)

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2 Table 2: Comparison of the eight different models specified in equations (1-8)

Rank	Model	Generalized linear model				Linear regression model				
		AICc	$\Delta_i$	$w_i$	$E_{8,i}$	Model	AICc	$\Delta_i$	$w_i$	$E_{8,i}$
1	8	505.1	0.0	0.9998	1	8	104.9	0	0.999	1
2	7	524.2	19.1	<0.0001	13805	3	120.3	15.4	0.0005	2211
3	3	524.8	19.7	<0.0001	19376	7	121.8	16.9	0.0002	4781
4	5	524.9	19.8	<0.0001	19641	5	121.9	17.1	0.0002	5083
5	6	527.5	22.42	<0.0001	73783	6	123.9	19.1	<0.0001	13940
6	4	527.5	22.43	<0.0001	74275	4	125.7	20.8	<0.0001	33247
7	2	538.5	33.4	<0.0001	>100000	2	129.7	24.8	<0.0001	>100000
8	1	541.3	36.2	<0.0001	>100000	1	130.7	25.8	<0.0001	>100000

3 Models were ranked according to increasing AICc, i.e., the higher AICc the less parsimonious the  
4 model and the worse the fit in relation to the number of variables employed. AICc: Bias-corrected  
5 Akaike Information Criterion;  $\Delta_i$ : Difference in AICc to the best model (models with  $\Delta > 15-20$  must  
6 be judged to be implausible);  $w_i$ : probability of model  $i$  being the Kullback-Leibler best model;  $E_{8,i}$ :  
7 evidence ratio between model 8 (the best model) and model  $i$

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11 Table 3 – Results of the full generalized linear models fitted to the original dataset with missing variables  
 12 removed (intercept not reported)

Variables	Full generalized linear model (N=31)		Full linear regression model (N=31)	
	Coefficient Estimate (SE)	p-value	Coefficient Estimate (SE)	p-value
log(test-standardized cases [%])	0.812 (0.143)	$7.6 \times 10^{-6}$ *	0.824 (0.174)	$7.9 \times 10^{-5}$ *
GRSI	0.015 (0.011)	0.192	0.012 (0.014)	0.393
Flu vaccination rate [%]	0.026 (0.005)	$8.6 \times 10^{-6}$ *	0.024 (0.006)	0.00026*
Vitamin D status (sufficient vs. deficient)	-0.777 (0.256)	0.0057*	--0.798 (0.311)	0.017
log( population density [ $\text{km}^{-2}$ ])	0.142 (0.097)	0.156	0.161 (0.117)	0.182
Gross domestic product	$2.8 \times 10^{-5}$ ( $7.3 \times 10^{-6}$ )	0.00064*	$3.1 \times 10^{-5}$ ( $8.8 \times 10^{-6}$ )	0.0016*
<b>Model quality</b>				
KL-R <sup>2</sup>	0.781		0.762	
Adjusted KL-R <sup>2</sup>	0.726		0.703	

13 GRSI: Government Response Stringency Index; SE: Standard error; \*p<0.01 (statistically  
 14 significant); \*p<0.01 (statistically significant)

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